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'Trans free vanaspati from palmstearin and
vegetable oils by interesterification process'

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Description

The present invention relates to improvements in edible fats, in particular to improved hardstocks for use in spreads having a low content of saturated fat and to spreads containing these hardstocks.

A vast number of edible fats are available for use in the manufacture of spreads and other fat containing foodstuffs. The rheological properties of these fats span a very wide range from low melting liquid oils to high melting fats with considerable variation in the shape and slope of melting and crystallization curves. In edible spreads it is desirable to use a liquid oil as the bulk of the fat, both for reasons of economy and nutrition. Consequently, the so-called "hardstocks" are needed to provide the structural component of spreads in the form of a fat in a crystalline phase at room temperature.

It is well accepted that the saturated fat content of the diet should be rather low. As hardstocks by necessity contain saturated fats, the quantity of hardstock in a product should be minimised. There is a need for hardstocks which function well at very low levels without a bad influence on the 'oral melt' of the product. A bad oral melt is a particular problem when the hardstock contains very high levels of saturated fatty acids as TAG's and in particular when the hardstock contains tripalmitic and tristearic TAG's.

A commonly used hardstock comprises a hydrogenated and interesterified blend of a palm oil (rich in C-16 and C-18 saturated fatty acids) with a so-called lauric fat (rich in C-12 saturated fatty acids). The best known examples of the lauric fats include coconut and palm kernel fats although a number of other tropical palm oils fall into this classification.

"BAILEY's industrial oil and fat products" Volume 2 fourth edition at page 159 describes how short chain fatty acids (C6-C14) improve the melting properties of spreads, while long chain fatty acids (C20-C22) provide stiffening power. BAILEY's goes on to describe how it is well known that both these properties can be combined using interesterified oil blends such as coconut/palm blends (as mentioned above), and coconut/hardened rape blends.

It must be appreciated that at least two different types of rapeseed oils are known. Originally, only so-called "high erucic acid" rapeseed or "HEAR" oil was known. Developments in crop breeding led to the production of so-called "low erucic acid rape", or "LEAR" oils. It is this second oil which is used in edible products.

FR 2570388 (GERSCHEL) discloses a fat composition for the manufacture of margarine produced by the interesterification of a hydrogenated vegetable oil including a low erucic acid rapeseed oil (colza) with a source of palmitic acid, such as palm fractions and technical tripalmitin. This is used as a hardstock ('mixed with fluid vegetable oil') in margarine manufacture. Looking at the GERSCHEL specification it can be appreciated that the rape seed oil is only partially hydrogenated and because it is low erucic acid rape oil it is rich in C-18 fatty acids. Although these products have good oral melting properties, with 20% hardstock levels, partially hydrogenated fats contain both *cis* and *trans* monosaturates, and these products are not acceptable to consumers, who wish to minimise their intake of *trans* fatty acids.

Edible fats are also subject to considerable variation in price. In particular, the lauric fats and other so-called "tropical" fats are cheap at present but may become more expensive in the future whereas liquid oils obtainable from more temperate crops are also relatively cheap but likely to remain so. A further problem with 'tropical fats' is that they are rich in C-16 fatty acids and is believed by some medical authorities that the intake of C-16 saturated fatty acids should be reduced. Most other edible oils and fats contain low levels of C-16 fatty acids and in particular edible liquid vegetable oils are rich in the nutritionally desirable C-18 unsaturated fatty acids.

Faced with the problems of tropical fats, workers have developed the techniques of "directed interesterification" to enable liquid vegetable oils to be used in spread hardstock production. However, directed interesterification is an expensive alternative to the use of these tropical fats.

EP-A-0 196 780 (Kao Corporation) discloses a fat bloom inhibitor for chocolate comprising a mixed acid triglyceride with 15-70% w.w. of C20-C24 saturated fatty acid residues and 20-60% w.w. of unsaturated C16-C22 fatty acid residues.

Synthesis example 5 discloses an interesterified blend of:

- (a) 33% fully hydrogenated high erucic rape seed oil
- (B) 17%? fully hydrogenated palm oil and
- (c) 50% unhydrogenated soybean oil

which rearranged oil as dissolved in acetone and fractionated. The liquid fraction was the desired inhibitor. It will be clear that the problem addressed here is totally different from that of the present invention. As to the interesterified fat blend disclosed it is clear that: (a) contains 46.9% of behenic acid (C22) and 5.5% of arachic acid (C20) totalling 52.4 of C20-C24 saturated fatty acids and the sum of palmitic and stearic acids is $43.6 + 3.9 = 47.5\%$; (b) contains 0.6% C20 and $43.6 + 54.2 = 97.8\%$ of C16 + C18 and (c) contains

10.4 + 4.1 = 14.5% of C16 + C18. The mixture consequently contains about 17.5% of C20-C24 and the amount of C16 + C18 = 0.33 x 45.2 + 0.17 x 97.8 + 0.50 x 14.5 = 14.9 + 16.6 + 7.3 = 38.8%. According to Table 1 the liquid fraction obtained therefrom contained 15.9 + 29.8 = 45.7% of C16 + C18, ipso facto the solid fraction must have been lower in C16 + C18 than the liquid fraction and even lower than the crude interesterified blend. Claim 1 of the present invention requires at least 50% of C16 + C18. Consequently Synthesis Example 5 is not even an accidental anticipation!

GB-A-1 205 729 (The Procter and Gamble Co) discloses plastic shortenings containing a combination (physical mixture) of hardstocks. For the most acceptable shortenings it is desirable that the solid glycerides crystallize and remain in the beta-prime crystalline form. Therefore the highest level possible of beta-prime phase stability is aimed at. For this reason certain mixtures of beta-prime tending hardstocks are provided. Hardstock (1) amounts to 5-50% of a C14-26 beta-prime phase-tending hardstock containing at least 20% (preferably 40%-60%) C20-22 fatty acids, whereas hardstock (2) amounts to 50-95% of a C14-18 beta-prime phase-tending hardstock containing at least 20% of C16-18 fatty acids, rather at least 20% C16. Preferably there is 10-30% of hardstock (1) and 90-70% of hardstock (2) present in the mixture of hardstocks used according to this specification in an amount of 4-15% in a plastic shortening.

There is at least one important difference between the hardstock used in accordance with this citation namely the there is no mention of the interesterification required according to the present invention which interesterification reshuffles the fatty acid residues over the hydroxyl groups of the glycerol molecule resulting in a random distribution (molecular rearrangement). Such a randomized fatty acid composition is e.g. expected to give one melting point (range) whereas the physical mixture of two hardstocks disclosed is expected to show two distinct and separate melting points in view of their different fatty acid compositions in the glycerides. Consequently an important difference in crystallizing behaviour is to be expected. Sandiness in margarine may a consequence.

We have now determined that hardstocks containing C16 (Palmitic - "P") or C18 (Stearic - "S") and C22 (Behenic - "B") saturated fatty acids combine structuring properties with a good oral melt when present at relatively low levels. In particular, our hardstocks comprise mixed triglycerides of P and/or S, and B fatty acid residues, in particular B2P and B2S TAGs. This reduces the level of use of tropical fats (normally present as P2M TAGs) and avoids use of directed interesterification. Moreover, these products contain low and preferably almost zero levels of trans fatty acids as the fats present are substantially fully hydrogenated.

Accordingly a first aspect of the present invention provides a fat blend, for use as a hardstock in an edible emulsion spread, obtainable by interesterification of a fat rich in behenic acid together with a fat rich in palmitic and/or stearic acid such that in the interesterified mixture the behenic acid content is at least 10% and the sum of palmitic and stearic acid content is at least 50%.

Typically, such hardstock fats may be obtained by interesterification of 10%-90% (by weight) wet fractionated palm stearin (80% palmitic acid) with 90%-10% (by weight) fully hydrogenated high erucic rapeseed oil. The preferred ratio of components is 40% of the behenic acid source and 60% of the palmitic and/or stearic acid source. For convenience it is noted that so called "wet fractionated palm stearin" comprises some 80% by weight of palmitic acid whereas high erucic acid rapeseed oil hardened to a slip-melting point of around 70 °C comprises around 48% behenic acid, the remainder being mainly C-18 fatty acids.

Typically, the hardstock and therefore the interesterified fat blend comprises around 27% behenic, 35% palmitic and 35% stearic acids.

Without wishing to be bound by theory, it is believed that the novel hardstocks with saturated C16/C22 + C18/C22 (palmitic/behenic) fatty acids (length ratio: 0.73) are in their behaviour as hardstocks similar to the known hardstocks with saturated C12/C16 (lauric/palmitic) fatty acids (length ratio: 0.75). The ratio for the C12/C22 (lauric/behenic) fatty acids, as mentioned above, is 0.55. The corresponding ratio for C18/C22 (stearic/behenic) fatty acids is 0.82. It is also expected that other randomised components of the fat blend will play some part in the behaviour of the blend as a hardstock.

Suitable sources of palmitic acid are fractions of palm oil, particularly wet or dry fractionated palm stearins and technical tripalmitin. The use of other sources of palmitic acid (such as genetically engineered plants) is not excluded.

Suitable sources of stearic acid are fully hydrogenated sunflower oil, safflower oil, soybean oil, corn oil, olive oil, and low erucic rapeseed oil.

Suitable sources of behenic acid are fully hardened high erucic acid rapeseed oil, fish oils and mustard seed oil. These fully hardened oils have a low content of trans fatty acids. Low erucic acid rapeseed oil has a low behenic acid content.

While the method of obtaining the fats disclosed herein is principally a chemical engineering method, the use of techniques of genetic engineering is not excluded. In particular, it is envisaged a medium erucic acid rapeseed oil rich in stearic and behenic acids randomised after hydrogenation to saturation would produce the hardstocks of the present invention. Such an oil can be produced by blending of low and high erucic acid rapeseed oils. This allows the preparation of a "one oil spread" with both hardstock and liquid-oil fat phase components derived from rapeseed fats.

Spreads including the present hardstock are included within the scope of the present invention, whether comprising rapeseed fats alone or not.

It is envisaged that as in the examples given hereafter the liquid oil component of the fat phase will generally be a liquid vegetable oil such as sunflower, safflower, groundnut or soybean oil, all of which contain relatively low levels of saturated fatty acids.

While the invention is described hereafter with particular reference to fat-continuous spreads comprising high levels of fat (around 80%) the hardstock of the present invention can find embodiment in spreads comprising 60%, 40% or even lower levels of fat. The production of water continuous spreads is not to be excluded. Spreads containing upwards of 60% fat can be used for frying and can further comprise suitable anti-spattering agents but it is suggested that spreads with lower fat levels, particularly 40%wt fat and lower are not used for this purpose. One or more of the fatty components of the oil phase may comprise a fat replacer such as a sucrose fatty acid ester. One or more of the components of the water-phase may comprise a fat replacer such as a finely dispersed protein.

In order that the present invention may be further understood it will be explained hereafter with reference to the following illustrative examples.

EXAMPLES:

A margarine was prepared with the following formulation (all percentages being given in wt%):

Hard fat component	4.90%
Sunflower oil	76.80%
Monoglyceride	0.10%
Lecithin	0.10%
Beta carotene	trace
Total fat phase	82.00%

Water	17.83%
Skim milk powder	0.12%
K sorbate	0.05%
Citric acid	(to pH 4.6)
Total water phase	18.00%

The hard fat (hardstock) component used at 6% on fat phase in both cases was either:

[A]: an interesterified blend (according to the present invention) of 40%wt wet fractionated palm stearin, and, 60%wt of fully hardened high erucic rapeseed oil with a slip melting point of 70 °C, or,

[B]: a simple mixture (as a comparative example) of 40%wt wet fractionated palm stearin, and, 60%wt of fully hardened rapeseed oil with a slip melting point of 70 °C.

EXAMPLE 1:

The spread according to both fat phase formulations, [A] and [B], was prepared according to the following process. Temperature measurements given are for the products made with the [A] fat phase:

All components of the spread were mixed together to form a premix at a temperature of 55 °C.

The premix was pumped at rate of 4.5 kg/h into a first "VOTATOR" A-unit rotating at 800 rpm with a cooling jacket temperature of -8 °C, an annulus of 3 mm spacing and a residence time of 15 seconds. The process stream outlet temperature for this unit was 12.5 °C and the solids content was measured at 4.2% by NMR.

From the first A-unit the process stream flowed to a second "VOTATOR" A-unit rotating at 800 rpm with a cooling jacket temperature of -3°C , an annulus of 3 mm spacing and a residence time of 15 seconds. The process stream outlet temperature for this unit was 4.8°C and the solids content was measured at 5.0% by NMR.

From the second A-unit the process stream was pumped to a "VOTATOR" C-unit rotating at 250 rpm with a cooling jacket temperature of $+13^{\circ}\text{C}$, a volume of 100 ml and a residence time of 120 seconds. The process stream outlet temperature for this unit was 8.5°C and the solids content was measured at 4.7% by NMR.

The product was filled into tubs.

The following physical properties were measured for the product according to the invention [A] and the comparative example [B]:

Sample	A	B
Solids content at stated temperature (%):		
N-10	7.4	6.1
N-20	5.4	4.7
N-30	3.9	3.7
N-35	2.5	2.2
Stevens value for consistency at temperature:		
S-05	47	20
S-10	44	19
S-15	47	19
S-20	32	19
Hardness value at stated temperature (calculated from Stevens value):		
C-05	210	95
C-10	195	90
C-15	210	90
C-20	150	90

It can be seen that the products according to the invention exhibit much improved consistency over the control products.

Whereas the consistency and hardness of the products of the invention are high at lower temperatures and remain quite high at room temperature, products made without the interesterified hardstock but otherwise comprising the same components are too soft for use as spreads at a range of temperatures.

Claims

1. Fat blend suitable for use as a hardstock in an edible emulsion spread, obtainable by interesterification of a fat rich in behenic acid together with a fat rich in palmitic and/or stearic acid such that in the interesterified mixture the behenic acid content is at least 8% preferably at least 10% and the sum of palmitic and stearic acid content is at least 50%.
2. Fat blend according to claim 1 obtainable by interesterification of 10%-90% (by weight) palm stearin with a fat rich in behenic acid.
3. Fat blend according to claim 1 obtainable by interesterification of 90%-10% (by weight) fully hydrogenated high erucic rapeseed oil with a fat rich in palmitic or stearic acid.
4. Fat blend according to claim 1 wherein ratio of components is about 40% of the behenic acid source and about 60% of the palmitic and/or stearic acid source.
5. Fat blend according to claim 1 wherein the fat rich in palmitic or stearic acid comprises more than 60% by weight of palmitic acid.

6. Fat blend according to claim 1 wherein the fat rich in behenic acid comprises more than 30% behenic acid.
7. Fat blend according to claim 1 wherein the fat rich in palmitic or stearic acid comprises wet or dry
5 fractionated palm stearins, technical tripalmitin, fully hydrogenated selected from sunflower oil, safflower oil, soybean oil, corn oil, olive oil, and low erucic rapeseed oil, and/or mixtures of the same.
8. Fat blend according to claim 1 wherein the fat rich in behenic acid comprises fully hardened high
10 erucic acid rapeseed oil, fish oils, mustard seed oil or mixtures of the same
9. Fat blend according to claim 1 comprising less than 10%wt, preferably less than 5% and most preferably zero trans fatty acids.
10. Edible water-in-oil emulsion spread comprising a fat blend according to claim 1 and a liquid vegetable
15 oil in a proportion such that the overall level of saturated fats present is less than 15%.

Patentansprüche

1. Fettgemisch, das zur Verwendung als hochschmelzende Fettkomponente (Hartfettkomponente oder
20 Hartfett) in einem eßbaren Emulsionsbrotaufstrich geeignet ist, erhältlich durch Interesterifizieren eines an Behensäure reichen Fettes zusammen mit einem an Palmitin- und/oder Stearinsäure reichen Fett, so daß in der interesterifizierten Mischung der Behensäuregehalt mindestens 8 %, vorzugsweise mindestens 10 %, beträgt und die Summe des Palmitin- und Stearinsäuregehaltes mindestens 50 % beträgt.
2. Fettgemisch nach Anspruch 1, erhältlich durch Interesterifizieren von 10 bis 90 Gew.-% Palmstearin mit einem an Behensäure reichen Fett.
3. Fettgemisch nach Anspruch 1, erhältlich durch Interesterifizieren von 90 bis 10 Gew.-% vollständig
30 hydriertem Rapsöl mit hohem Erucasäuregehalt mit einem an Palmitin- oder Stearinsäure reichen Fett.
4. Fettgemisch nach Anspruch 1, bei dem das Verhältnis der Komponenten etwa 40 % Behensäurequelle und etwa 60 % Palmitin- und/oder Stearinsäurequelle beträgt.
- 35 5. Fettgemisch nach Anspruch 1, bei dem das an Palmitin- oder Stearinsäure reiche Fett mehr als 60 Gew.-% Palmitinsäure umfaßt.
6. Fettgemisch nach Anspruch 1, bei dem das an Behensäure reiche Fett mehr als 30 % Behensäure umfaßt.
- 40 7. Fettgemisch nach Anspruch 1, bei dem das an Palmitin- oder Stearinsäure reiche Fett naß oder trocken fraktionierte Palmstearine, technisches Tripalmitin, vollständig hydriertes pflanzliches Öl, ausgewählt aus Sonnenblumenöl, Distelöl, Sojaöl, Maisöl, Olivenöl und Rapsöl mit niedrigem Erucasäuregehalt, und/oder Mischungen derselben umfaßt.
- 45 8. Fettgemisch nach Anspruch 1, bei dem das an Behensäure reiche Fett vollständig gehärtetes Rapsöl mit hohem Erucasäuregehalt, Fischöle, Senföl oder Mischungen derselben umfaßt.
9. Fettgemisch nach Anspruch 1, das weniger als 10 Gew.-%, vorzugsweise weniger als 5 Gew.-% und
50 insbesondere keine trans-Fettsäuren umfaßt.
10. Eßbarer Wasser-in-Öl-Emulsionsbrotaufstrich, umfassend ein Fettgemisch nach Anspruch 1 und ein flüssiges pflanzliches Öl in einem solchen Verhältnis, daß die Gesamtkonzentration der vorliegenden gesättigten Fette weniger als 15 % beträgt.

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Revendications

1. Un mélange gras approprié pour être utilisé en tant que matière solide dans une pâte à tartiner d'émulsion comestible, pouvant être obtenu par interestérification d'une graisse riche en acide béhénique avec une graisse riche en acide palmitique et/ou stéarique de façon telle que dans le mélange interestérifié la teneur en acide béhénique est au moins égale à 8 %, de préférence au moins égale à 10 %, et la somme de la teneur en acide palmitique et stéarique est au moins égale à 50 %.
2. Mélange gras selon la Revendication 1, pouvant être obtenu par interestérification de 10 % à 90 % (en masse) de stéarine de palme avec une graisse riche en acide béhénique.
3. Mélange gras selon la Revendication 1, pouvant être obtenu par interestérification de 90 % à 10 % (en masse) d'huile de colza fortement érucique entièrement hydrogénée avec une graisse riche en acide palmitique ou stéarique.
4. Mélange gras selon la Revendication 1, dans lequel le rapport des composants est environ égal à 40 % pour la source d'acide béhénique et environ égal à 60 % pour la source acide palmitique et/ou stéarique.
5. Mélange gras selon la Revendication 1, dans lequel la graisse riche en acide palmitique ou stéarique comprend plus de 60 % en masse d'acide palmitique.
6. Mélange gras selon la Revendication 1, dans lequel la graisse riche en acide béhénique comprend plus de 30 % d'acide béhénique.
7. Mélange gras selon la Revendication 1, dans lequel la graisse riche en acide palmitique ou stéarique comprend des stéarines de palme fractionnées sèches ou humides, de la tripalmitine technique, de l'huile entièrement hydrogénée sélectionnée parmi de l'huile de toumesol, de l'huile de carthame, de l'huile de soja, de l'huile de maïs, de l'huile d'olive et de l'huile de colza faiblement érucique, et/ou des mélanges de celles-ci.
8. Mélange gras selon la Revendication 1, dans lequel la graisse riche en acide béhénique comprend de l'huile de colza à acide fortement érucique, des huiles de poisson, de l'huile de graine de moutarde entièrement durcies ou des mélanges de celles-ci.
9. Mélange gras selon la Revendication 1, comprenant moins de 10 % en masse, de préférence moins de 5 %, et de façon plus préférentielle zéro acides gras trans.
10. Pâte à tartiner comestible d'émulsion eau-dans-huile comprenant un mélange gras selon la Revendication 1 et une huile végétale liquide dans une proportion telle que la quantité totale de graisses saturées présentes est inférieure à 15 %.